

## General

#### Title

Brain and central nervous system (CNS) cancer: proportion of patients with malignant glioma (with enhancing component on pre-operative imaging) undergoing surgical resection who receive MRI within 3 days (72 hours) of surgical resection.

## Source(s)

NHS Scotland, Scottish Cancer Taskforce. Brain and central nervous system cancer clinical quality performance indicators. Edinburgh (Scotland): Healthcare Improvement Scotland; 2016 Jan. 39 p. [24 references]

#### Measure Domain

#### Primary Measure Domain

Clinical Quality Measures: Process

# Secondary Measure Domain

Does not apply to this measure

# **Brief Abstract**

## Description

This measure is used to assess the proportion of patients with malignant glioma (with enhancing component on pre-operative imaging) undergoing surgical resection who receive magnetic resonance imaging (MRI) within 3 days (72 hours) of surgical resection.

Note from the National Quality Measures Clearinghouse: This measure is part of the Cancer Quality Performance Indicators (QPIs) collection. For more information, including a complete list of QPI measure sets, please visit the Healthcare Improvement Scotland Web site

#### Rationale

Post-operative imaging:

Provides a measurement of surgical performance;

Helps to determine if further treatment is required; Helps determine what further treatment might be appropriate; Estimates residual tumour to help target radiotherapy when needed; and Helps to assess prognosis.

Imaging should be carried out within 72 hours to enable reliable assessment of the extent of the resection (van den Bent et al., 2009; Ulmer et al., 2006; Cairncross et al., 1985; Sato et al., 1997; Smith et al., 2005). Magnetic resonance imaging (MRI) is the preferred imaging method for patients with glioma.

After this time period, changes in the tumour resection bed confound estimation. Delaying assessment until these changes settle is inappropriate as regrowth of high-grade tumours can occur rapidly and also post-operative treatments such as radiotherapy and chemotherapy are normally instituted rapidly which could further affect the images.

#### Evidence for Rationale

Cairncross JG, Pexman JH, Rathbone MP, DelMaestro RF. Postoperative contrast enhancement in patients with brain tumor. Ann Neurol. 1985 Jun;17(6):570-2. PubMed

NHS Scotland, Scottish Cancer Taskforce. Brain and central nervous system cancer clinical quality performance indicators. Edinburgh (Scotland): Healthcare Improvement Scotland; 2016 Jan. 39 p. [24 references]

Sato N, Bronen RA, Sze G, Kawamura Y, Coughlin W, Putman CM, Spencer DD. Postoperative changes in the brain: MR imaging findings in patients without neoplasms. Radiology. 1997 Sep;204(3):839-46. PubMed

Smith JS, Cha S, Mayo MC, McDermott MW, Parsa AT, Chang SM, Dillon WP, Berger MS. Serial diffusion-weighted magnetic resonance imaging in cases of glioma: distinguishing tumor recurrence from postresection injury. J Neurosurg. 2005 Sep;103(3):428-38. PubMed

Ulmer S, Braga TA, Barker FG 2nd, Lev MH, Gonzalez RG, Henson JW. Clinical and radiographic features of peritumoral infarction following resection of glioblastoma. Neurology. 2006 Nov 14;67(9):1668-70. PubMed

van den Bent MJ, Vogelbaum MA, Wen PY, Macdonald DR, Chang SM. End point assessment in gliomas: novel treatments limit usefulness of classical Macdonald's Criteria. J Clin Oncol. 2009 Jun 20;27(18):2905-8. PubMed

# Primary Health Components

Brain/central nervous system (CNS) cancer; malignant glioma; surgical resection; magnetic resonance imaging (MRI)

# **Denominator Description**

All patients with malignant glioma (with enhancing component on pre-operative imaging), undergoing surgical resection (see the related "Denominator Inclusions/Exclusions" field)

# **Numerator Description**

Number of patients with malignant glioma (with enhancing component on pre-operative imaging), undergoing surgical resection receiving magnetic resonance imaging (MRI) within 3 days (72 hours) of surgical resection (see the related "Numerator Inclusions/Exclusions" field)

# Evidence Supporting the Measure

## Type of Evidence Supporting the Criterion of Quality for the Measure

A clinical practice guideline or other peer-reviewed synthesis of the clinical research evidence

A formal consensus procedure, involving experts in relevant clinical, methodological, public health and organizational sciences

One or more research studies published in a National Library of Medicine (NLM) indexed, peer-reviewed journal

## Additional Information Supporting Need for the Measure

Unspecified

## **Extent of Measure Testing**

The collection of data is piloted on a small number of patient records using a paper data collection form produced by the Information Services Division (ISD). The aim is to identify any anomalies or difficulties with data collection prior to full implementation. At least one NHS board in each Regional Cancer Network participates in the pilot.

## Evidence for Extent of Measure Testing

NHS Scotland. National cancer quality performance indicators: overview of development process. Edinburgh (Scotland): NHS Scotland; 2012 Dec. 7 p.

# State of Use of the Measure

#### State of Use

Current routine use

#### **Current Use**

not defined yet

# Application of the Measure in its Current Use

# Measurement Setting

Ambulatory Procedure/Imaging Center

Hospital Outpatient

## Professionals Involved in Delivery of Health Services

not defined yet

## Least Aggregated Level of Services Delivery Addressed

Single Health Care Delivery or Public Health Organizations

#### Statement of Acceptable Minimum Sample Size

Unspecified

## Target Population Age

Unspecified

## **Target Population Gender**

Either male or female

# National Strategy for Quality Improvement in Health Care

# National Quality Strategy Aim

Better Care

## National Quality Strategy Priority

Prevention and Treatment of Leading Causes of Mortality

# Institute of Medicine (IOM) National Health Care Quality Report Categories

#### IOM Care Need

Getting Better

Living with Illness

#### **IOM Domain**

**Timeliness** 

## Data Collection for the Measure

## Case Finding Period

Unspecified

## **Denominator Sampling Frame**

Patients associated with provider

#### Denominator (Index) Event or Characteristic

Clinical Condition

Diagnostic Evaluation

Therapeutic Intervention

#### **Denominator Time Window**

not defined yet

## Denominator Inclusions/Exclusions

Inclusions

All patients with malignant glioma\* (with enhancing component on pre-operative imaging), undergoing surgical resection

\*Malignant gliomas include:

Glioblastoma multiforme (GBM) and its variants (e.g., gliosarcoma)
Anaplastic astrocytoma (AA)
Anaplastic oligodendrogliomas
Mixed tumours (e.g., oligoastrocytoma, glioblastoma with oligodenroglial component)
High-grade ependymoma

#### Exclusions

Patients unable to undergo a magnetic resonance imaging (MRI) scan, e.g.:

Pacemaker or other MRI incompatible implanted device

Cerebral aneurysm clip

Contraindication to intravenous contrast medium

Patients who refuse MRI

Patients undergoing biopsy only

Note: Where it is not possible to image with MRI an attempt should be made to image with computerised tomography (CT).

# Exclusions/Exceptions

not defined yet

#### Numerator Inclusions/Exclusions

#### Inclusions

Number of patients with malignant glioma (with enhancing component on pre-operative imaging), undergoing surgical resection receiving magnetic resonance imaging (MRI) within 3 days (72 hours) of surgical resection

#### Exclusions

Patients unable to undergo a MRI scan, e.g.:

Pacemaker or other MRI incompatible implanted device

Cerebral aneurysm clip

Contraindication to intravenous contrast medium

Patients who refuse MRI

Patients undergoing biopsy only

Note: Where it is not possible to image with MRI an attempt should be made to image with computerised tomography (CT).

## Numerator Search Strategy

Fixed time period or point in time

#### **Data Source**

Electronic health/medical record

Paper medical record

## Type of Health State

Does not apply to this measure

## Instruments Used and/or Associated with the Measure

Unspecified

# Computation of the Measure

# Measure Specifies Disaggregation

Does not apply to this measure

## Scoring

Rate/Proportion

# Interpretation of Score

Desired value is a higher score

# Allowance for Patient or Population Factors

not defined yet

#### Standard of Comparison

not defined yet

#### Prescriptive Standard

Target: 90%

The tolerance within this target is designed to account for situations where patients are deemed unfit to attend for imaging within the stated timeframe.

#### **Evidence for Prescriptive Standard**

NHS Scotland, Scottish Cancer Taskforce. Brain and central nervous system cancer clinical quality performance indicators. Edinburgh (Scotland): Healthcare Improvement Scotland; 2016 Jan. 39 p. [24 references]

# **Identifying Information**

#### **Original Title**

QPI 7 - early post-operative imaging.

#### Measure Collection Name

Cancer Quality Performance Indicators (QPIs)

#### Measure Set Name

Brain and Central Nervous System Cancer

#### Submitter

NHS Scotland - National Government Agency [Non-U.S.]

Scottish Cancer Taskforce - National Government Agency [Non-U.S.]

## Developer

NHS Scotland - National Government Agency [Non-U.S.]

Scottish Cancer Taskforce - National Government Agency [Non-U.S.]

# Funding Source(s)

## Composition of the Group that Developed the Measure

Brain/Central Nervous System (CNS) Cancer QPI Development Group

#### Financial Disclosures/Other Potential Conflicts of Interest

Unspecified

## Adaptation

This measure was not adapted from another source.

## Date of Most Current Version in NQMC

2016 Jan

#### Measure Maintenance

The Cancer Quality Performance Indicators (QPIs) will be kept under regular review and be responsive to changes in clinical practice and emerging evidence.

## Date of Next Anticipated Revision

2017 Aug

#### Measure Status

This is the current release of the measure.

# Measure Availability

Source document available from the Healthcare Improvement Scotland Web site
·
For more information, contact the Healthcare Improvement Scotland at Gyle Square, 1 South Gyle
Crescent, Edinburgh, Scotland EH12 9EB; Phone: 0131 623 4300; E-mail: comments.his@nhs.net; Web
site: www.healthcareimprovementscotland.org/

# Companion Documents

The following is available:

HS Scotland. National cancer quality performance indicators: overview of development process.	
dinburgh (Scotland): NHS Scotland; 2012 Dec. 7 p. This document is available from the Healtho	are
mprovement Scotland Web site	

This NQMC summary was completed by ECRI Institute on May 4, 2017. The information was verified by the measure developer on May 23, 2017.

#### Copyright Statement

No copyright restrictions apply.

#### **Production**

## Source(s)

NHS Scotland, Scottish Cancer Taskforce. Brain and central nervous system cancer clinical quality performance indicators. Edinburgh (Scotland): Healthcare Improvement Scotland; 2016 Jan. 39 p. [24 references]

#### Disclaimer

#### NQMC Disclaimer

The National Quality Measures Clearinghouseâ, (NQMC) does not develop, produce, approve, or endorse the measures represented on this site.

All measures summarized by NQMC and hosted on our site are produced under the auspices of medical specialty societies, relevant professional associations, public and private organizations, other government agencies, health care organizations or plans, individuals, and similar entities.

Measures represented on the NQMC Web site are submitted by measure developers, and are screened solely to determine that they meet the NQMC Inclusion Criteria.

NQMC, AHRQ, and its contractor ECRI Institute make no warranties concerning the content or its reliability and/or validity of the quality measures and related materials represented on this site. Moreover, the views and opinions of developers or authors of measures represented on this site do not necessarily state or reflect those of NQMC, AHRQ, or its contractor, ECRI Institute, and inclusion or hosting of measures in NQMC may not be used for advertising or commercial endorsement purposes.

Readers with questions regarding measure content are directed to contact the measure developer.